

## MICROARRAYS AND THEIR MANUFACTURE

### CROSS REFERENCE TO RELATED APPLICATIONS

5 This application is a continuation-in-part of Provisional Patent Application Serial No. 60/146,653 filed July 30, 1999, the content of which is incorporated herein in entirety. The content of Disclosure Document Nos. 286,465 filed July 11, 1991; 418,726 filed May 9, 1997; 444,028 filed August 31, 1998; 445,769 filed October 5, 1998; and 458852 filed 14 July 1999 are  
10 hereby incorporated herein by reference.

### FIELD OF THE INVENTION

The present invention relates to microarrays containing bioreactive molecules, their uses and methods for their manufacture. The arrays are  
15 constructed by sectioning bundles of tubules or rods containing unique reactants to produce large numbers of identical arrays.

### BACKGROUND OF THE INVENTION

Microarrays are known in the art and are commercially available from a  
20 number of sources. Microarrays have been used for a number of analytical purposes, typically in the biological sciences. An array is essentially a two-dimensional sheet where different portions or cells of the sheet have different biomolecule elements, such as, nucleic acids or peptides, bound thereto. Microarrays are similar in principle to other solid phase arrays except that  
25 assays involving such microarrays are performed on a smaller scale, allowing many assays to be performed in parallel. For example, the reactive biomolecules bound to a chip.

Biochemical molecules on microarrays have been synthesized directly at or on a particular cell on the microarray, or preformed molecules have been attached to particular cells of the microarray by chemical coupling, adsorption or other means. The number of different cells and therefore the number of different biochemical molecules being tested simultaneously on one or more microarrays can range into the thousands. Commercial microarray plate readers typically measure fluorescence in each cell and can provide data on thousands of reactions simultaneously thereby saving time and labor. A representative example of the dozens of patents in this field is U.S. Pat. No. 5,545,531.

Currently two dimensional arrays of macromolecules are made either by depositing small aliquots on flat surfaces under conditions which allow the macromolecules to bind or be bound to the surface, or the macromolecules may be synthesized on the surface using light-activated or other reactions. Previous methods also include using printing techniques to produce such arrays. Some methods for producing arrays have been described in "Gene-Expression Micro-Arrays: A New Tool for Genomics", Shalon, D., in *Functional Genomics: Drug Discovery from Gene to Screen*, IBC Library Series, Gilbert, S.R. & Savage, L.M., eds., International Business Communications, Inc., Southboro, MA, 1997, pp 2.3.1.-2.3.8; "DNA Probe Arrays: Accessing Genetic Diversity", Lipshutz, R.J., in *Functional Genomics: Drug Discovery from Gene to Screen*, IBC Library Series, Gilbert, S.R. & Savage, L.M., eds., International Business Communications, Inc., Southboro, MA, 1997, pp 2.4.1.-2.4.16; "Applications of High-Throughput Cloning of Secreted Proteins and High-Density Oligonucleotide Arrays to Functional Genomics", Langer-Safer, P.R., in *Functional Genomics: Drug Discovery from Gene to Screen*, IBC Library Series, Gilbert, S.R. & Savage, L.M., International Business Communications,

Inc., Southboro, MA, 1997, pp 2.5.13; Jordan, B.R., "Large-scale expression measurement by hybridization methods: from high-densities to "DNA chips"", J. Biochem. (Tokyo) 124: 251-8, 1998; Hacia, J.G., Brody, L.C. & Collins, F.S., "Applications of DNA chips for genomic analysis", Mol. Psychiatry 3: 483-92, 1998; and Southern, E.M., "DNA chips: Analyzing sequence by hybridization to oligonucleotides on a large scale", Trends in Genetics 12: 110-5, 1996.

Regardless of the technique, each microarray is individually and separately made, typically is used only once and cannot be individually precalibrated and evaluated in advance. Hence one depends on the reproducibility of the production system to produce error-free arrays. These factors have contributed to the high cost of currently produced biochips or microarrays, and have discouraged application of this technology to routine clinical use.

For scanning arrays, charged coupled device (CCD) cameras are widely used. The cost of these has declined steadily, with suitable cameras and software now widely available. However, in one proposed variation, an array is located at the ends of a bundle of optical fibers with the nucleic acid or antibody/antigen attached to the end of the optical fiber. Detection of fluorescence may then be performed through the optical fiber, see U.S. Pat. No. 5,837,196.

Fiber optical arrays are routinely produced in which glass or plastic fibers are arrayed in parallel in such a manner that all remain parallel, and an optical image may be transmitted through the array. Parallel arrays may also be made of hollow glass fibers, and the array sectioned normal to the axis of the fibers to produce channel plates used to amplify optical images. Such devices are used for night vision and other optical signal amplification equipment.

Channel plates have been adapted to the detection of binding reactions (US Pat. No. 5,843,767) with the individual holes being filled after sectioning of the channel plate bundle, and discrete and separate proteins or nucleic acids being immobilized in separate groups of holes.

5       Hollow porous fibers have been widely used for dialysis of biological samples, in kidney dialyzers and for water purification. Methods for aligning them in parallel arrays, for impregnating the volume between them with plastic, and for cutting the ends of such arrays have been described (see, for example, U.S. Pat. No. 4,289,623).

10       Immobilized enzymes have been prepared in fiber form from an emulsion as disclosed in Italy Pat. No. 836,462. Antibodies and antigens have been incorporated into solid phase fibers as disclosed in U.S. Pat. No. 4,031,201. A large number of other different immobilization techniques have been used and are well known in the fields of solid phase immunoassays, nucleic acid hybridization assays and immobilized enzymes, see, for example, 15       Hermanson, Greg, T. Bioconjugate Techniques. Academic Press, New York. 1995, 785 pp; Hermanson, G.T., Mallia, A.K. & Smith, P.K. Immobilized Affinity Ligand Techniques. Academic Press, New York, 1992, 454 pp; and Avidin-Biotin Chemistry: A Handbook. D. Savage, G. Mattson, S. Desai, G. 20       Nielander, S. Morgansen & E. Conklin, Pierce Chemical Company, Rockford IL, 1992, 467 pp.

Scanners and CCD cameras have been described to detect and quantitate changes in fluorescence or adsorbance and are suitable for existing biochips. These, together with suitable software, are commercially available.

25       Currently available biochips include only one class of immobilized reactant, and perform only one class of reactions. For many types of clinical and other analyses there is a need for chips which can incorporate in one chip

reactants immobilized in different ways.

## SUMMARY OF THE INVENTION

The present invention relates to a method for producing rods or tubules,  
5 each containing a different entrapped biological agent of interest, for arranging  
and keeping the rods or tubules in parallel bundles, for impregnating or  
embedding the bundles with a sectionable adhesive material, for checking that  
all elements of the bundle maintain a constant arrangement or pattern  
10 throughout the length of the bundle after impregnation, for sectioning the  
bundle to produce large numbers of identical arrays or chips, and for  
performing a variety of different quantitative biochemical analyses on  
individual arrays or chips based on enzymatic or immunochemical activities  
under conditions yielding fluorescence or optical absorbence signals, for  
15 acquiring images of these signals which are electronically processed and  
compared to produce clinically and experimentally useful data.

In one aspect, the invention relates to long filaments or tubes that  
contain, are coated with, or have an agent of interest embedded therein, and  
methods for their manufacture.

The invention also relates to methods for arranging the fibers to form  
20 bundles in which the position of each fiber relative to all others is retained  
throughout the bundle length.

The invention further relates to means and methods for attaching or  
gluing all of the fibers together over their entire length.

In a related aspect, the invention relates to the preparation of microarrays  
25 wherein the elongated filaments or tubes are bundled together and cut  
transversely many times at short intervals to form microarrays and a microarray  
so prepared.

A further aspect of the invention is the inclusion of markers which are either integral with the tubes or fibers or are contained in the media contained in hollow fibers which allow them to be distinguished along the entire length.

5 An additional aspect of the invention includes means for illuminating fibers individually at one end of a bundle, and identifying the other end by photoelectric means to confirm the integrity of the fiber arrangement.

In another aspect, the present invention relates to forming a fiber containing an agent of interest, or means for immobilizing one or a class of agents of interest.

10 In an additional aspect, the invention relates to means for embedding or attaching whole or fragments of cells, tissues or infectious agents to fibers of tubules in such a manner that these are exposed on the cut end of each fiber of tubule.

15 In another aspect of the invention, the array consists of tubules containing gel or other polymerizing materials that adhere to the tubing walls.

In a further aspect of the invention, agents of interest are attached to the polymerizing or suspending medium in the lumen of the small tubes.

20 In yet another aspect of the invention, the agents of interest are attached to particles that are suspended in a polymerizing medium, which suspension is used to fill tubules used to make array bundles and arrays.

This invention further relates to a method for the large scale production of identical flat two-dimensional arrays of immobilized nucleic acid-based agents for use in nucleic acid sequencing in the analysis of complex mixtures of ribonucleic acids (RNAs) and deoxyribonucleic acids (DNAs), and in the  
25 detection and quantitation of other analytes including proteins, polysaccharides, organic polymers and low molecular mass analytes, by sectioning long bundles of fibers or tubes.

In a related aspect, the invention relates to exploiting microarrays for mass screening of large numbers of samples from one to a large number of agents of interest.

In a further related aspect the invention relates to the development of sets of tests on different chips done in sequence, which reduces the cost, delay, and inconvenience of diagnosing human diseases, while providing complex data ordinarily obtained by time-consuming sequential batteries of conventional tests.

In still another aspect, the invention relates to the fabrication of identical arrays which are sufficiently inexpensive to allow several identical arrays to be mounted on the same slide or test strip, and cross compared for quality control purposes.

In a still further aspect, the invention relates to the incorporation of a non-fluorescent dye or other light absorbing material in the substance of the array to control the depth to which light used to excite fluorescence penetrates the array, thereby controlling the depth to which fluorescence analytes are detected, and insuring that fluorescent analytes which diffuse too deeply into the contents of the cells and therefore do not diffuse out are not detected.

In another aspect, the invention relates to methods for determining that tubules are completely full of support media, and lack voids or air bubbles.

In a further aspect, the invention relates to methods and apparatus for completely filling small tubes with a supporting medium using hydrostatic force or centrifugal force.

In an additional aspect, the invention relates to the reproducible manufacture of biochips for bioanalysis.

In a further aspect, the invention relates to the design and production of arrays, which are specifically designed to detect and diagnose a specific

disease.

In yet another aspect, the present invention relates to multi-welled plates and methods for their manufacture.

5 In yet a further aspect, the invention relates to increasing the dynamic range of multiple-parallel assays by providing means for making serial measurements of fluorescence or absorbance over time, and for determining the rate of change of fluorescence or absorbance in each element of the array over time.

10 It is an additional aspect of this invention to produce biochips which are inexpensive and sufficiently standardized to allow more than one to be used for each analysis, and for controls and standards to be routinely run simultaneously in parallel. For added quality assurance, sections from different portions of the bundle or different ends may be used. One way of sectioning from different portions of the bundle is to cut or bend the bundle in the middle and align the  
15 two halves to form a single larger bundle thereby producing a section where each fiber is represented twice.

In a further aspect, this invention relates to the production of chips in which the array elements or cells may differ from one another in the composition of the tubes, supporting medium, immobilization surface, or the  
20 class of agent of interest may be different in different cells.

In an additional aspect, the invention relates to the production of chips in which different types of reactions may be carried out at the surface of each cell of the array, with the reactions including immunological, enzymatic or hybridization reactions.

25 The invention further relates to the development of multiple parallel chip-based methods involving continuously increasing temperature such that temperature sensitive reactions may be carried out at physiological



temperatures, followed by an increase in temperature to allow hybridization reactions to occur.

In a still further aspect, the invention relates to preparing libraries of compounds with each fiber containing one of the compounds. The array may be used to simultaneously screen all of the compounds for a particular chemical or biological activity.

### BRIEF DESCRIPTION OF THE DRAWINGS

Figure 1 is a schematic of intermediate products in the process for producing microarrays.

Figure 2 is a schematic of an individual tubule containing beads with immobilized ligands embedded in a gel.

Figure 3 is a schematic of an individual tubule containing a gel with ligands attached to the gel.

Figure 4 is a schematic of an array with ligands attached to the inner walls of cells, and with means for closing off one surface of the array to form a set of microwells.

Figure 5 is a schematic of a means for insuring that all fibers are maintained in their correct pattern before the bundle is sliced.

Figure 6 is a schematic of means for identifying arrays.

Figure 7 is a schematic for scanning an array.

Figure 8 displays an alternative way of forming a fiber bundle.

### DETAILED DESCRIPTION OF THE INVENTION

The terms "binding component", "molecule of interest", "agent of interest", "ligand" or "receptor" may be any of a large number of different molecules, biological cells or aggregates, and the terms are used

interchangeably. Each binding component is immobilized at a cell, site or element of the array and binds to an analyte being detected. Therefore, the location of an element or cell containing a particular binding component determines what analyte will be bound. Proteins, polypeptides; peptides, nucleic acids (oligonucleotides and polynucleotides), antibodies, ligands, polysaccharides, microorganisms, receptors, antibiotics, test compounds (particularly those produced by combinatorial chemistry), bacteria, viruses, or plant and animal cells and organelles or fractions of each may each be a binding component if immobilized on the chip. Each, in turn, may also be considered as analytes if they bind to a binding component on a chip.

When a molecule of interest has a high molecular weight, it is referred to as a “macromolecule”. In terms of some biopolymers, the high molecular weight refers to greater than 100 amino acids or nucleotides or sugar molecules long.

The term “bind” includes any physical attachment or close association, which may be permanent or temporary. Generally, an interaction of hydrogen bonding, hydrophobic forces, van der Waals forces etc. facilitates physical attachment between the molecule of interest and the analyte being measuring. The “binding” interaction may be brief as in the situation where binding causes a chemical reaction to occur. This is typical when the binding component is an enzyme and the analyte is a substrate for the enzyme. Reactions resulting from contact between the binding agent and the analyte are also within the definition of binding for the purposes of the present invention.

The term “cells”, “sites” or “elements” in this application refers to a unit component of an array identified by a unique address and these generally differ from other cells, sites or elements by their content as well as location. Biological cells are referred to by their type, e.g. microorganisms, animal and

plant cells.

The term "fibers" includes both filaments and hollow capillaries. Filaments or rods may be solid strands of monolithic, porous, or composite forms, or aggregate forms. Pluralities, typically a large number, of fibers are bound adjacent to each other in ribbons or bundles to form a "fiber bundle." The cross-section of the fibers may be of any shape, such as round, triangular, square, rectangular or polygonal.

The term "particle" includes a large number of insoluble materials of any configuration, including spherical, thread-like, brush-like and many irregular shapes. Particles are frequently porous with regular or random channels inside them. Examples include silica, cellulose, Sepharose beads, polystyrene (solid, porous and derivitized) beads, controlled-pore glass, gel beads, sols, biological cells, viruses, subcellular particles, etc. Even certain high molecular weight materials, such as, polymers and complexes, may serve as immobilizing structures that would constitute a "particle".

The terms "arrays" and "microarrays" are used somewhat interchangeably differing only in their general size. The present invention involves the same methods for making and using either. Each array typically contains many cells (typically 100-1,000,000+) wherein each cell is at a known location and contains a specific component of interest. Each array therefore contains numerous different components of interest.

The present invention makes microarrays or "chips" by sectioning bundles of small plastic rods, fibers, tubes or tubules containing immobilized binding component, including biological molecules and entities such as nucleic acid fragments, antigens, antibodies, proteins, peptides, carbohydrates, ligands, receptors, drug targets, biological cells or their subfractions, infectious agents or subfractions of them, drugs, toxic agents, or natural products. Embedding

media may be, in the present invention, polymerized or solidified in small tubes, or may be cast into rods or sheets.

The tubes may be of material such as glass, metal, ceramic or plastic.

The immobilized binding components, e.g. nucleic acids, proteins, cells, etc.,

5 may be coated on the inside of the microtubes, contained in a gel in the microtubes, or attached to small particles or beads which fill the tubes. The particles or beads may be a component of a gelling material or can be separate components such as latex beads made of a variety of synthetic plastics (polystyrene, etc.). When the individual fibers are solid rods or filaments, the  
10 agent of interest is incorporated into the plastic before the filament is cast, extruded or pulled through a die. Each section cut constitutes a microarray for use in various binding assays.

Various aspects of the invention are illustrated Figures 1-7. General principles are illustrated in Figure 1 where rod or tube 1 incorporates an agent  
15 of interest. The rods or tubes may be bonded into a flat parallel array 2, and multiple flat arrays are then bonded into the multiple-parallel bundle 3.

Alternatively the bundle 3 may be constructed in one step from a series of rods 1. The end of bundle 3 is cut or sectioned to yield the final array 4 which contains one small section 5 of each rod or tube in the entire bundle. By making  
20 a long bundle 3, and cutting very small sections 4, a very large number of identical arrays or chips are formed. For example, if bundle 3 is a meter long, and the sections are 10 microns thick, 100,000 identical chips may be produced.

For the present purposes, in the case of hollow glass fibers, such as those  
25 in channel plates, the hollow fibers may be filled with gels or particles including immobilized reactants, and the entire bundle sawed into arrays.

The rods or tubules comprising the sectioned bundle fall into at least

eight classes, with subdivisions of each.

A **first class** is composed of solid rods or filaments with the immobilized binding component being part of the composition of the rod or filament. The agent of interest in the present invention may comprise a very broad range of

5 chemicals, complexes, tissues, biological cells or fractions thereof. Nucleic acids, proteins, which may have been modified or are coated with detergents to make them more soluble in organic solvents, and a wide range of organic compounds can be incorporated into polymerizing mixtures such as those used to produce plastics. Oligonucleotides and nucleic acids are soluble in  
10 methylene chloride, for example, and hence may be included in acrylics during polymerization.

A number of polymerizing embedding agents have been developed for histological and histochemical studies, some of which are listed in Table 1, together with data on their composition, curing temperature, solvent used and  
15 viscosity.

TABLE I

RESIN*	TYPE	CURE TEMP.	SOLVENT	VISCOSITY
Durcupan	-	40°C	Water	Medium
Nanoplast	Melamine	60°C	Water	Low
Quetol 651	Epoxy	60°C	Water	Low
London Resin Gold	Acrylic, UV Curing	-25°C	Water, EtOH	Low
Lowicryl K4M Polar	Acrylic, UV Curing	-35°C	Water, EtOH	Low
Lowicryl Monostep K4Mpolar	Acrylic, UV Curing	-35°C	Water, EtOH	Low
Lowicryl K11 Polar	Acrylic UV Curing	-60°C	EtOH	Low
JB-4	GMA	RT	Water	Low
JB-4 Plus	Methacrylate	RT	Water	Low
ImmunoBed	GMA	RT	Water	Low
PolyFreeze	Polyol	-15°C	Water	Low

\*Available from Polysciences

- 5        **A second class** of fibers are not homogeneous and the polymerizing or gelling material may also contain solid structural elements such as filaments, branched elements etc., to further strengthen the gel or and may also provide attachment sites for the agent of interest. Thus the added components serve to strengthen the gel, and may provide attachment sites for inclusions including
- 10    dendrimer branched polynucleic acids, branched or crosslinked polymeric materials, metal or glass fibers. Threads, yarn-like and brush-like configurations of structural elements may be cast into the length of the fiber giving it strength and allowing the fiber to be more easily handled or dried. The structural elements may serve as the immobilizing component in the fiber
- 15    for a desired binding component.

Thus it is technically feasible to produce long fibers of acrylic or other plastics each containing a different agent of interest using currently available extrusion technology in the present invention. The cut end of the fibers may be briefly treated with dilute solvents to expose active groups.

5        **A third class** of fibers includes extruded or cast plastic which includes a second phase. This second phase may be in the form of, for example, hydrocarbon, aqueous or fluorocarbon microdroplets, particles of sugars or other water soluble materials, or inorganic particles such as calcium carbonate particles, which can be dissolved in dilute acid to reveal active groups. Brief  
10 exposure of the cut surface of a chip to a solvent will dissolve some of these inclusions, increasing the surface area of the support plastic containing the agents of interest.

Solid plastics can also be prepared which incorporate polystyrene latex or other plastic particles to which proteins or nucleic acids are attached.  
15 Conditions can be arranged such that the supporting plastic is eroded to a depth of a few microns to reveal active subparticle surfaces, and do not dissolve the supporting plastic latex beads. For example, proteins derivatized with fluorinated groups attach strongly to Teflon® microparticles. Such derivatized Teflon® particles in, for example, an acrylic plastic or other suitable embedding  
20 medium, can be partially exposed at the plastic surface by a dilute acrylic solvent, composed, for example, of methylene chloride and ethyl alcohol.

The beads to which agents of interest are attached may be porous gel beads used in chromatography such as Sephadex, Biogels and others, or solid beads such as are used in chromatography. A variety of methods for  
25 derivitizing these and for attaching proteins, nucleic acids and polysaccharides and small molecules thereto have been developed and are well known to those skilled in the arts. The construction of the tubules of this class is illustrated in

Figure 2 where tubule 6 is comprised of tube 7 containing gel 8 which supports particles 9. An end view 10, and enlarged view 11 of the tubule shows exposed particles 12 at the cut end. Area 13 is shown additionally enlarged at 14 to illustrate the presence of immobilized reactants 15 on the surface of the exposed particles 12.

Note that all rods described can be cast with a string or thread through their center to increase strength, and to make the rods easier to handle.

**A fourth class** of fibers is prepared by sintering glass or plastic beads to form a porous material with a high surface to mass ratio. Such material is conventionally made from glass, polytetrafluoroethylene (PTFE) (Teflon®), Teflon®AF, polyethylene, polypropylene, can be manufactured from polystyrene and from a variety of other plastics. Glass can be stiff. Heat, pressure, or exposure to solvent vapors can sinter plastics. The sintered material can be derivatized in sheets or in cut rods. Polystyrene is the most convenient from the point of view of coupling agents of interest thereto. For polystyrene derivatization, methods that allow attachment of proteins by their amino groups, carboxyl groups, or sulfhydryl groups, have been described and widely used. Teflon® can be activated using solutions of metallic sodium in an organic solvent producing groups to which other plastics will adhere, and these may then be derivatized. Polyethylene and polystyrene can be activated by corona plasma discharge or by electron beam radiation. However the most advantageous approach appears to be to make sintered composites of polystyrene and polyethylene. Nylon beads can also be sintered and derivatized. Other sintered materials are known or are under development, many of which will find application here.

Molecules of interest may be attached to the solid materials either before or after sintering. For attachment of the ligands, the rods may be soaked in



tubes containing the substance to be attached or the rods may be coiled up inside a hollow bowl centrifuge rotor having the general configuration of a zonal rotor (see Anderson, N.G., Natl. Cancer Inst. Monograph No. 21), but which may be centrifugally drained. The solution of the substance to be  
5 attached is then centrifuged first into the sintered mass, and then out of it, followed by washing as necessary. The sintered rods may then be dried, coated with a suitable adhesive, assembled into a bundle and sectioned.

Alternatively, the beads with nucleic acid sequences attached may be extruded under pressure to form rods that may then be sintered together. The  
10 assembled tubes may be held together with a variety of cements or polymerizeable plastics. The outsides of the tubes may be altered or treated so that cements or polymerizeable plastics will adhere to them.

**A fifth class** of fibers is comprised of hollow impermeable tubules typically formed from plastics including, but not limited to polyethylene,  
15 polypropylene, Teflon®, or polyvinyl chloride, and is completely filled with a gel or other polymerizing material to which agents of interest are attached directly. The external surfaces of the tubes may be chemically or physically modified to accept adhesives used to bind the bundled tubes together. The internal surface may also be modified so that the gel or polymerizing mixture  
20 introduced into the tubes will adhere, preferably by covalent attachment. Acrylamide derivatives may be linked to the wall to make an acrylamide gel adhere, while gelatin, agar, or agarose derivatives may be similarly attached to link with their respective gels. Methods for linking proteins and nucleic acids to linear acrylamide, gelatin and agarose are well known, and these derivitized  
25 molecules can be incorporated into the gels used for casting. Acrylamide can be made to gel at room temperature either chemically or using photoactivation, while low temperature-gelling Sepharose is readily available. Gelatin sets

slowly, and at temperatures below ambient. The polymers used to fill the tubes are typically homogeneous, but may contain agents of interest which become attached to the polymerizing medium. Examples include covalent attachment of proteins to short acrylamide chains that become incorporated into acrylamide gels and proteins covalently linked to gelatin. Thus gels are available or can be produced which contain labile biomolecules without exposing them to denaturing temperatures. The structure of tubes of this class is illustrated in Figure 3 where tube 16 is filled with a cross-linked gel 17 to which are attached agents of interest 18. A side view 19 and end view 21 of a sectioned tube illustrates the availability of immobilized agents 21.

Arrays prepared using hollow fibers may have the interior of the fibers coated with biomolecules either covalently or in suitable polymer coatings, or in gels before the array is assembled. Alternatively, a bundled array may be positioned so that individual hollow fibers may be filled with biopolymers in solutions which gel, prior to sectioning.

**A sixth class** of fibers or tubes includes empty impermeable tubes with molecules of interest attached to the inner surface, but otherwise empty or made empty. As illustrated in Figure 4, the sectioned chip 22 is comprised of sectioned plastic tubes 23 embedded in supporting plastic 24, with the agent of interest 25 attached to the inner walls of the tubes, leaving the center 26 open. The result 27, seen in side section, has sectioned plastic tube 28, immobilized agent 29, yielding open holes 30, and all held together by supporting material 31. The chips may be considered as ultramicrotiter plates, and may be used for flow through analysis based on immobilized affinity ligand techniques (Hermanson et al., Immobilized Affinity Ligand Techniques, Academic Press, 1992, p 407), for polymerase chain reaction (PCR) amplification of immobilized oligonucleotides, or for other reactions as described in US Pat.

No. 5,843,767. When the tubes are made of Teflon® with the internal or external surfaces treated to become hydrophilic, the cut ends will remain hydrophobic. When a hydrophilic test solution is spread across the surface of the chip, it tends to flow into the holes in self-controlling volumetric amounts, and, if the total amount of fluid is properly controlled, tend not to affect adjacent cells. The upper and lower surfaces can then be sealed with a suitable adhesive tape, and the whole subject to reactions, for example, for polymerase chain reaction amplification of DNA. Alternatively the sandwiched structure including chip of Figure 4 may employ two pieces of material such as glass or quartz to seal the ends of the tubes, creating microchambers. Changes in fluorescence or in optical absorbance may be detected in each tubular element through the transparent end windows, and the reaction followed colorimetrically or fluorometrically.

The inside surfaces of the small tube described may be modified chemically to allow attachment of nucleic acids, proteins or other molecules either directly or through linkers known per se. The molecules attach, thus increasing the number of reactive sites inside the tube. Since DNA and RNA are conventionally synthesized on small polystyrene beads, the most direct approach to a nucleic acid array is to synthesize oligonucleotides on small polystyrene beads, with different batches of beads having different sequences attached, and to then fill small polyethylene, polypropylene, polystyrene or other plastic, metal or ceramic tubes with the beads, packing them down to completely fill them. The beads may be kept in place by carefully heating them to sinter or residual latex is added to the tubes and dried in place with air pumped through the tube.

**A seventh class** of tubes or fibers includes tubules with permeable walls. Methods and procedures for producing hollow selectively permeable fibers for

use in kidney dialysis machines and for molecular weight fractionation have been extensively developed (US Pat. No. 4,289,623, US Pat. No. 3,976,576) and are in wide current use. Procedures for embedding such fibers in solid sectionable plastics have also been developed and are routinely used to attach  
5 the fibers to tubing at the dialyzer ends.

Permeable hollow fibers may be used in the present invention in two ways. In the first, the fibers are filled with reactant-carrying gels while already embedded in plastic. By carefully splaying out the fibers going into the cast portion, each tube can be selectively filled as previously described. This  
10 technique offers the advantage of producing small arrays quickly, and of developing new assays without having to go through all of the steps required to produce separate hollow fibers, fill them with reactants, arrange them in arrays, and infiltrate them with the supporting plastic.

The second method of use involves filling the hollow fibers before they  
15 are embedded in plastic. Techniques have been developed for controlling the wall permeability of permeable tubes. This allows the influx and outflux of monomers and gelling agents during gelation to be controlled, and for dialyzable agents to be removed after gelling. For example, acrylamide gels may be produced from acrylamide and bisacrylamide by cross-linking with  
20 ultraviolet light in the presence of riboflavin. This catalyst, which would interfere with subsequent fluorescence measurements, can be removed by dialysis through the tubing wall after polymerization.

One preferred gelling material, polyacrylamide, may be polymerized by ultraviolet (UV) light using a catalyst such as riboflavin. This technique is  
25 preferred where the specific binding component is heat sensitive or sensitive to other chemicals. The riboflavin may then be removed by washing or elution.

Permeable supporting tubing also allows the gel inside a tube to be

infiltrated with substances that render with the reactants more stable, increase the physical strength of the gel, and facilitate sectioning. For example sugars such as lactose or trehalose may be introduced to stabilize proteins, and to add solids to the gels to assist in sectioning. These additives may be partially removed from the exposed surface of the chip during use to make buried reactive groups available. Additives diffusing into the gels may also be used to increase the strength and volume of a gel after it has been dried.

Also when a particle containing the ligand or receptor is embedded in a fiber, the embedding medium may be soluble or meltable so as to be removable after the microarray is formed. By removing the embedding medium, more active sites on the particle are exposed for binding. This variation is particularly preferred when the particle is actually microfibers or microbrushes of microfilaments having the immobilized ligands or receptors thereon similar to the cross-linked polymers of (17) in Figure 3.

Once the tubes are filled with their respective gels and reagents, the outside of the tubes are cleaned, may be treated with reagents to increase the adherence of the infiltrating supporting plastic, and then bundled to produce the product for sectioning.

An eighth class of tubes or fibers includes those synthesized by cleaving from a larger block, preferably a disk. The fiber material containing the molecule of interest is first cast as a disk and then a long fiber is peeled from the circumference of a rotating disk. This technology is essentially the same as a smaller version of producing wood veneers where the veneer is peeled from a rotating log. This technique has certain space and handling advantages over a long thin fiber. Such a disk also is more easily stored, particularly when active components therein require maintenance under certain conditions, e.g. freezing, submergence in buffer, in the dark, etc.

Arrays or parallel fibers may be attached together by many techniques. Preferred ones are by vapor sintering or by heat solvent vapor being introduced.

The vapor is allowed to interact with the array for a specified period of time, and is then removed by re-evacuation. In heat sintering, the array is placed  
5 under lateral compression and the array heated to the softening point of the plastic. Another means is the use of low melting point metals, such as gallium. By low melting point is meant temperatures at or about physiologic temperature of the binding component

A variety of histological embedding media have been developed that  
10 which preserves nucleic acids and antigens in a reactive form. These include, among others, Durcupan, Nanoplast, Quetrol 651 which may be cured by very mild heating, JB-4, Immunobed which may be polymerized at room temperature, and the water soluble acrylic polymers London Resin Gold and Lowicryl which are polymerized at below freezing temperatures by ultraviolet  
15 light (all are available from Polysciences Inc.). Conventional embedding media use solvents and waxes, and the waxes must be at least partially removed before analysis.

Embedding and sectioning methods are therefore available which allow  
*in situ* hybridization to identify and localize specific DNA or RNA sequences,  
20 and amplification of specific sequences by the polymerase chain reaction and other nucleic acid amplification techniques (LCR, RCA, SDA, etc).

The method of embedding is one which preserves the desired characteristic or characteristics of the binding component in a cell. Thus, if antibodies were immobilized in a cell and it is the antigen-binding specificity  
25 of the antibody that is desired, the immobilization method will be one which retains the antigen-binding ability of the antibodies. The method and means of attaching the fibers to form the array are also ones which retain the antigen-

binding ability of the antibodies.

Similarly, if the cells contain candidate molecules for binding to a hormone receptor, the immobilizing and attaching method and means are those which retain the configuration of the candidate molecules which allows

5 recognition and binding by the hormone receptor.

In addition, many protein or carbohydrate antigens may be detected using immunological reagents. Detection is generally by incorporation of a fluorescent dye into the analyte or into the second layer of a sandwich assay, or by coupling an enzyme to an analyte or a second or third layer of a sandwich  
10 assay which produces an insoluble dye which may be fluorescent.

Some solid phase surfaces may be used directly to immobilize reactants, others must be modified to allow such additions. Antibodies will adhere to clean polystyrene surfaces, as will many other proteins (Van Oss, C.J., & Singer, J.M. The binding of immune globulins and other proteins by  
15 polystyrene latex particles. J. Reticuloendothelial Society 3: 29040, 1966.) Polystyrene, either in the form of microtiter plates or beads, have been modified to bind nucleic acids, proteins, and polysaccharides using techniques that are well known. Teflon® surfaces will tenaciously bind proteins or other macromolecules that have been suitably fluorinated (U.S. Pat. No. 5,270,193),  
20 and will also bind fluorinated surfactants, which may render the surface hydrophilic, or positively or negatively charged. Glass, including controlled pore glass, may be modified to allow covalent attachment of antibodies, antigens or nucleic acids. Plastic or glass surfaces may be modified non-specifically using corona plasma discharge or electron beam radiation and may  
25 then be coated with a variety of coatings or adhesives to which macromolecules may be attached. More specific covalent attachment of proteins, nucleic acids or carbohydrates may be achieved by a variety of modifications which attach

reactive groups to polystyrene or acrylic surfaces, which groups, with or without extending linkers, will then couple under mild conditions to the biopolymers.

A variety of chromatographic media have also been adapted to support immobilized bioreactants. These include soft gel beads, generally composed of acrylamide, agarose, Sepharose, which may be chemically cross-linked, and less compressible beads designed for high-pressure chromatography. Chief among natural products useful as an immobilization support is cellulose that is readily available in powdered form. These supports may be chemically modified to allow covalent bioreactant attachment, or may be purchased in modified form ready for attachment.

Long DNA or RNA molecules may be immobilized by being polymerized in a gel and are retained purely by physical entanglement. An example of this is the retention of DNA in agar or acrylamide gels. In addition, proteins or nucleic acids may be covalently linked to long polymer chains so that, when embedded in a gel, they are prevented from diffusing out, but are still available for reaction with soluble reactants. Examples of this include the attachment of proteins or nucleic acids to polyethylene glycol (so-called PEGylation), or to linear acrylamide chains.

In addition to methods by which a receptor or molecule of interest is immobilized and used to bind an analyte, general methods exist for arranging to immobilized members of a class of reactants. For example, protein A or protein G may be immobilized and used to subsequently bind specific immunoglobulins which in turn will bind specific analytes. A more general approach is built around the strong and specific reaction between other ligands and receptors such as avidin and biotin. Avidin may be immobilized on a solid support or attached to a gel and used to bind antibodies or other reactants to



which biotin has been covalently linked. This allows the production of surfaces to which a very wide variety of reactants can be readily and quickly attached (see Savage et al., *Avidin-Biotin Chemistry: A Handbook*. Pierce Chemical Company, 1992).

5 A wide variety of methods have been developed to detect reactions between immobilized molecules of interest and soluble reactants. These differ chiefly in the mechanism employed to produce a signal, and in the number of different reagents which must be sandwiched together directly or indirectly to produce that signal. These include fluorescence (including delayed  
10 fluorescence) with the fluorescent tag covalently attached to the analyte, fluorescence involving soluble dyes which bind to an analyte, and similar dyes whose fluorescence greatly increases after binding an analyte. The latter are chiefly used to detect nucleic acids. In more complex systems, including so-called sandwich assays, the end result is the immobilization in the detection  
15 complex of an enzyme that, in combination with a soluble substrate, produces a preferably insoluble dye that may be fluorescent. Alternatively, the detection complex attached to the bound analyte may include a dendritic molecule, including branching DNA, to which is attached many fluorescent dye molecules.

20 Methods for making dental floss having attached short transverse fibers to give a brush-like configuration have been described, and these may be modified to allow attachment of reactants. Patterns encoding identifying information on strands or fibers may be employed in the form of small linearly arranged dots. In the development of multifiber endoscopy arrays, methods for  
25 checking the array have been developed in which a light beam or raster image is introduced at one end of the fiber bundle in such a manner that the light sequentially illuminates each fiber. The pattern of emitted light exiting the

other end is then determined. If they are identical, no fiber is out of place.

The art of detecting bubbles or voids in liquid filled tubing is well known, and may depend on differences in refraction, or light absorption or fluorescence as measured along individual tubes. The art of using centrifugal force to fill short lengths of small tubing with viscous media is evident to those trained in the arts.

Microtomes for sectioning tissue blocks which may contain soft tissues in wax all the way to blocks including bone, are commercially available, as are a variety of techniques and arrangements for attaching sections to glass or plastic slides, for treating them automatically to remove some or all of the embedding media, and for systematically exposing the slides to a series of reagents.

Microtomes and other sectioning or cutting instruments capable of cutting assembled bundles of tubes into thin sections, and of maintaining their orientation after sectioning are known. In general, blade cutting is preferred to sawing to reduce contamination of binding components between cells of the microarray.

The sections (as microarray chips) may be attached directly to adhesive surfaces on flexible films or on solid surfaces, such as glass slides. It is also feasible to attach sections (the word "section" is used here in place of "chip") at intervals along a film strip, with others interleaved between them. Thus a set of about a dozen or more different sections may be placed in repeating order along the film, and the film then cut up to give one set. For sequencing studies, one DNA insert may be amplified, labeled, and its hybridization to a large set of sections examined.

By using a non-deformable bundle of fibers, one can cut or saw the bundle transversely thereby forming a large number of identical plates that are

perfectly realignable. This permits highly consistent and reproducible arrays. By using an easily detectable different material for one or more fibers, as a means for registering the microarray alignment, realignment is even easier.

Most immunochemical or competition assays depend on a signal  
5 produced by a reagent other than the analyte. However, methods for  
fluorescently labeling all proteins containing aliphatic amino groups in a  
complex mixture have been developed which are reproducible and quantitative.  
Of these CyDyes supplied by Amersham Life Sciences, and particularly, Cy2,  
Cy3 and Cy5 have proven most useful. When the components of such labeled  
10 mixtures are reacted with an array of immobilized antibodies, each specific  
antibody to one of the fluorescently labeled analytes, the presence of each of  
the specifically bound labeled analyte can be detected by fluorescence. This  
method can be further improved by exposing the bound antibody array to a  
solution containing known subsaturating quantities of each protein in a non-  
15 fluorescent form, washing the bound antibody array, and exposing it to a test  
mixture of labeled proteins, thus producing a multiple competition assay.

Any of the conventional binding assay formats involving an immobilized  
binding partner, known per se in hundreds of patents, may be used with the  
microarray systems of the present invention.

20 It is well known that several different clinical tests are often required to  
define a particular disease. These are often done serially, with one test or  
member of a battery of tests suggesting another, which in turn suggests a third  
test or group of tests, some of which are rarely done in local laboratories.  
Frequently one series of tests suggests one or more additional series to modify,  
25 or confirm a diagnosis. There is therefore a need for inexpensive chips for the  
performance of branching batteries of tests using methods that produce accurate  
numerical results in a machine readable form, which are stable over time, and

which are read by devices which can be compact and inexpensive relative to currently clinical analytical systems.

Many biochemical analyses require that the analytical procedure have wide dynamic range. Thus enzyme and immunochemical assays are often done by determining the course of a reaction over a period of time, or by doing the analyses on a series of dilutions. In addition, parallel analyses using standards and blanks (controls) are required and are universally included. Large numbers of standardized inexpensive biochips will be required to meet these needs.

These biochips may incorporate reactants of different classes that can, for example, detect and measure antigens, drugs, nucleic acids or other analytes simultaneously.

Arrays have numerous uses other than determining bioactive properties. Chemical interactions and reactions may be tested as well such as an array of different reactive chemicals being tested against a test substance or material to determine corrosion, electrochemical reaction or other interaction. This is particularly advantageous in the chemical formulations of plural substances such as in cosmetics, paints, lubricants, etc. Alternatively, one may assay for desirable interactions between the analyte and all of the molecules of interest in the array.

A general problem with use of gels for the immobilization of reactants has been that reactants, which may attach to gel-immobilized agents of interest require considerable time to diffuse into and out of the gel. Where the detection is by fluorescence, inclusion of a light absorbing dye into the gel limits detection to a region close to the surface. Inclusion of the ultraviolet light absorbing monomer 4-methacryloxy-2-hydroxybenzophenone (Polysciences, Inc.) in an acrylic embedding medium solves this problem.

When one wishes to enhance binding between analyte and binding

partners on the surface area of particles in a fiber of the microarray, one may etch the embedding matrix of each fiber, thereby exposing more of the surface area of particles in each fiber of the microarray.

When performing a binding assay, one may wish to encourage diffusion of the analyte into the microarray cell to increase ligand/receptor binding (sensitivity), to make the microarray more quantitatively reproducible and to enhance spectrophotometric detection if done by passing light through the microarray. To enhance diffusion through the microarray one may force the ligand through the microarray gel material. This may be done by mounting the microarray on a porous membrane and passing the ligand and or ligand solution through the microarray by hydrodynamic, electrophoretic or mechanical means. For example fluid may be flowed through the microarray by pressure difference on each side of the membrane. Fluid may also be drawn through by simply applying a stack of paper towels on the backside of the membrane to suck fluid through the microarray. As for electrophoretic means, a potential is applied across the microarray either across the entire microarray or using single point electrodes located on both sides of a single or group of cells of the microarray. Mechanical means may involve a pump of various configurations to mechanical push or pull fluid through the microarray by providing a pressure differential.

Using a porous membrane also has certain advantages in washing the microarray to achieve lower backgrounds. If porous particles or threadlike components are embedded within the fiber, sectioning through the porous particle or threadlike component may make it more porous and allow greater surface area contact to both reagents and washing. Etching of an embedding medium or capillary also increases porosity and exposure to the immobilized molecules of interest.

If a porous particle is sectioned, preferably twice, larger channels allowing more fluid passage may be present. Fibers with these sectioned particles may be mounted over permeable membrane supports or over holes in a solid base support. The result allows fluid to pass through the cells of the  
5 microarray.

By using the present invention, one avoids the difficulties of individually spotting each cell on a solid phase or forming a compound at each cell. The former technique is limited by the spill and ability to quantitatively measure small quantities of liquid. The later technique is limited by the types of  
10 different compounds that can be synthesized on the solid phase. Both prior art techniques are expensive and require elaborate automated equipment or tedious labor as each array is individually produced. By contrast, the present invention is technically simple and quick where the "batch" is in the thousands to millions of microarrays. The only individual effort required for each  
15 microarray is the step of cutting.

The agent of interest in the present invention may comprise a very broad range of chemicals, complexes, biological cells or fractions thereof. Nucleic acids, many proteins, proteins which have been modified or are coated with detergents such as sodium dodecyl sulfate are soluble in organic solvents and a  
20 wide range of organic compounds and thus can be incorporated into polymerizing mixtures such as those used to produce plastics. Hence it is technically feasible to produce long fibers of acrylic or other plastics each containing a different agent of interest using currently available extrusion technology for practice in the present invention.

25 Large numbers of different and potentially new active compounds may be simultaneously screened by immobilizing them in fibers, bundling, sectioning and forming a microarray. Peak fractions from separations, such as

plant extracts may be simultaneously collected and used to form a microarray. The microarrays may then be used in a large number of assay systems simultaneously, dramatically reducing the time and effort to screen all of the compounds present for whatever activity one chooses.

5            Particularly preferred are large numbers of proteins or peptides generated by mass techniques. Different fractions from a separation technique from a natural source provide a resource of many different proteins and peptides. A number of fractionation procedures are known to separate mixtures of many compounds. Different fractions or specific compositions may be used to form  
10 a single fiber. Two dimensional electrophoresis gels from serum and other tissue and natural sources produces tens of thousands of different proteins separated on the gel. Each may be individually removed (e.g. cut, eluted etc.) from the gel and used as the molecule of interest to form a single fiber. In such a method, with different bundles being formed from different samples, protein  
15 differences between different samples may be readily compared.

When the immobilized macromolecules are antibodies, the microarray may be used to diagnose a variety of protein-based anomalies. A labeled second antibody to the protein of interest may be used to further highlight the cell. In addition the array may be used to immobilize infectious agents which  
20 have been either previously stained or which are stained after immobilization. Thus, microbes from biological samples, e.g. serum or plasma, may be concentrated, stained with a fluorescent nucleic acid stain such as TOTO-1 or YOPRO-1, and then allowed to find their matching antibodies on the array. They may then be detected by scanning for fluorescence and identified by  
25 position.

It is equally a part of the present invention to immobilize microorganisms or other molecules of interest and use them to localize

antibodies from a patient's sera, and then discover the location of the latter using a fluorescent anti-human antibody, thus diagnosing a disease which elicited antibody production in the first place.

Arrays have been prepared using phage display with inserts from specific genes, using synthetic oligonucleotides, or (to a limited extent) using displayed antigens or antibodies. In the present application a population of peptide or antibody display phage may be used where each display phage is used to prepare a single fiber. In such an arrangement, the phage is large enough so that some portion of each surface molecule will remain embedded in the gel or plastic, while part of it will be exposed. The molecule of interest may be bound to the fiber per se, entrapped inside the matrix, or bound to a solid phase particle or tiny structure which is in or on the fiber.

The each fiber may contain a mixture of molecules of interest. For example, during chemical synthesis, a number of isomers are prepared. It is convenient to not separate the isomers before forming a fiber in some circumstances. Likewise, when fractionating a mixture, forming a fiber with a mixture of receptors may be acceptable as total and complete isolation is difficult and time consuming.

The embedding matrix for the fibers may be black, opaque or otherwise adsorbent to emitted signals of a label in order to reduce cross talk between the cells in the chip. Additionally, any adhesive between the fibers may contain the same adsorbent material to reduce background between cells of the microarray. Optionally, a specific layer of this material may be placed between the fibers before they form the bundle. When hollow fibers are used, the opaque material may be incorporated into the hollow fiber shell itself.

Arrays may have an entire set of antigens/antibodies etc. in the various cells along with controls to effectively screen blood samples for common blood



borne diseases before donated blood is provided for transfusion. Likewise, certain symptoms have a number of common causes that may be simultaneously screened for using arrays. For example, urinary tract infections are common and may be caused by a large number of different bacteria of varying sensitivity to various antibiotics. The simultaneous testing for numbers factors would save considerable time and expense.

In the course of using a chip of the instant invention, various known techniques and materials are used to reduce non-specific reaction. Thus, in the case of a protein-based assay, the non-specific sites on the chip contributed by the substance of the fiber or filament, the embedding material and essentially everything aside from the binding component of interest are reacted with a blocking agent, such as albumin or milk, so that the blocking agent will bind to those areas not containing the binding component which could react with an ligand, analyte, reporter molecule or whatever would specifically bind to the binding component, as known in the art.

Arrays may have two or more identical cells made from different fibers but containing identical binding agents. This provides an internal quality assurance check for the array. Additionally, it is preferred for some of the cells to provide different concentrations of the binding component for quantitative measurement of an analyte. These provide internal standards for the microarray for both qualitative detection and quantitative detection. For example, a series of cells may contain different concentrations of an antibiotic in their gels. When a sample microorganism is contacted with the cells and allowed to incubate, the absence of growth in one cell and the presence of growth in another cell provide an approximate minimal inhibitory concentration. The same can be done for determining minimal bacteriocidal concentrations when stained with a vital dye such as trypan blue or fluorescein

acetate. Since a microarray may contain thousands of cells, one can simultaneously determine the antibiotic sensitivity to numerous antibiotics simultaneously. Quantitative determination of other biological activities with either ligand or receptor immobilized in the gel may be used.

5 Essentially the same fiber may be used multiple times in the same microarray. This provides an internal quality control check and improves confidence in the binding assay. This also provides additional quantitative measurements if such an assay is performed to improve precision. Blank fibers, fibers with no molecule of interest bound thereto, provide a good  
10 negative control and should be used in every microarray.

Long filaments, capillaries or coaxial two-material filaments are arranged in parallel and then sintered or adhesively bonded to each other to form bundles which are preferably resistant to deformation, and in which each strand or capillary is continuous from one to the other. The positional arrangement of  
15 fibers or capillaries should remain the same throughout the bundle. Filaments composed of two different types of material in coaxial formation may be used. The core material is made of a material, which can be dissolved, and the cladding being resistant to the same dissolving conditions. For example, strong alkali is capable of dissolving certain types of glass but not others. The  
20 dissolving step may occur before or more preferably after sectioning depending on the materials present.

Alternatively, the cladding may be dissolvable and the core resistant leaving isolated "islands" on a microarray attached to a backing sheet. In either situation, the space left by the dissolving step may remain empty or be filled  
25 with a diverse material. Partial dissolving to yield a porous material is also part of the present invention. Porous materials have increased surface area, which is particularly desirable for binding assays.

Particles, especially porous beads, may also be “chemically sintered” to form a filament, sheet or inside of a capillary. This technique may also be used to adhere different fibers together. One such way is to first bind a molecule of interest to the particle. A blocking agent may be added to block any remaining  
5 active sites or adsorption areas on the particle. If not already done, the beads are packed in a tube or the hollow fiber. A chemically reactive compound which crosslinks or couples either the blocking agent and/or the molecule of interest and/or unreacted sites on the beads is then added and at the locations where the beads touch, chemical adhesion results. The tube or hollow fiber  
10 may remain in place or be removed. Since the molecules of interest in the internal pores of the beads are not touching, they are not significantly altered. Alternatively, the pores of the beads may be filled with a hydrophilic solution and held by capillary action while the spaces between the beds are filled with a hydrophobic adhesive or setting liquid.

15 A representative example of chemical sintering is to adsorb Protein G on porous beads and then adding a gelatin blocking agent. The resulting beads are filled in a 1mm plastic tube and then a protein crosslinking agent added, e.g. carbodiimide. After the reaction is complete, unreacted reagents are washed free and then any suitable antibody of interest is added thereto to bind to  
20 Protein G to form a fiber suitable for bundling and cleaving to make a microarray.

Alternatively, the surfaces of the particles may be biotinylated first and avidin may be used as the crosslinking agent. One may even use avidin labeled antibodies instead of adsorbing Protein G to the beads. Another alternative is  
25 to use relatively large porous beads and use an adhesive or embedding medium to fill the spaces between the beads. When the fiber is sectioned, the beads are so large that they are cleaved, thereby opening up the inside of the beads for the

bound molecules of interest to be exposed. Hollow beads or microballoons may be used in lieu of porous beads as molecules of interest encapsulated therein will be exposed upon cleavage of the bead. The concept is the same as sectioning a tissue or embedded cell to expose and visualize intracellular

features.

Additionally, one may use two different sets of beads: set one is porous and has the receptor/receptor binding substance bound thereto, and the second set is coated with highly reactive material or modified with a reactive group which will bind to the first set of beads or coating thereon. A tube is first filled with both beads in dry form, the tube shaken, and then fluid is pumped therethrough permitting a reaction to occur thereby forming a solid fiber of beads. Alternatively, if the first set of beads is quite large they may be added first (with or without fluid) and the second set added later so that they filter down through the spaces between the larger beads and react accordingly. The reaction between the beads may be through specific binding moieties or of a non-specific binding reaction to form a crosslinking of the beads into a sliceable solid. The second beads may be black to reduce stray light in the fluorescence detection.

After the fibers in the bundle are fused or otherwise adhered to each other in a fixed pattern, the bundle is cut transversely or at an angle into many thin disks and portions are optionally dissolved if desired. When hollow capillaries are used, the resulting disks may be used as channel plates for the amplification of optical images and light pipes. Regardless of whether rods or fibers are used, the thin disks may also be used as filters because of their remarkably uniform hole size.

Each fiber segment in the sectioned two-dimensional array would contain relatively large numbers of binding components, such as DNA, RNA,

or protein molecules. As a first step in the use of the final array, a solution, which can erode the plastic surface of the array very slowly, is washed over the surface. This is done at a rate which will remove any biopolymer molecules which become loose. The wash is then continued, grading into a solution which will not erode the plastic. The array may then be dried and stored until used, or may be used at once. To assist in exposing reactive agents of interest in the plastic, particles on the surface are dissolved, forming a solution, and exposing the molecules.

For clinical tests, regulatory approval of tests and systems and methods for making them is required. When chips are fabricated using photolithography and other technology derived from electronic chip making, the cost of individual chips is extraordinarily high, and the possibility of error when chips are individually made is very high. Since chips are individually made and used only once, quality control is difficult and there is no good way of proving that any given chip is satisfactory. The best that can be done is to test a large fraction of a batch at random. With the present invention, a very large number of sections can be made from one composite assembly, and adjacent sections intercompared as well as those some distance apart. Statistical analyses will be able to predict the rate of errors that may occur. However, of even greater importance is the fact that since the sections can be made in large numbers and quite cheaply, it will be feasible to run duplicate analysis on clinical samples, and to run confirmatory analysis when important diagnostic results are obtained. The present invention therefore makes feasible widespread and routine application of genetic analyses in the practice of medicine.

The key agent of interest components of the fibers is retained by the fiber by being immobilized therein. Immobilization may be accomplished by a number of techniques, known per se, such as entrapment in a matrix and

chemically coupled, perhaps through a linking moiety through an amino, hydroxy, sulfhydryl or carboxyl moiety. Chemically attaching the chemical to a monomer to be polymerized also effectively incorporates the component. Binding may also be accomplished by a number of affinity techniques such as protein A or protein G for antibody attachment, ligand/receptor pairs such as biotin-avidin, HIV-CD4, or through a ligand that has a receptor such as digoxigenin-antidigoxigenin. On the other hand, no specific attachment is required for situations where a gel or a non-gel, gelling matrix, such as wax, is used. Liquid wax or a gelling agent is simply mixed with the key component and cooled to form a solid fiber by casting or extruding.

Arrays need not be assembled in a single step. Flat arrays consisting of a set of tubes arranged side-by-side may be prepared first, and the end of the array sectioned and tested. The flat arrays can then be attached together with a suitable adhesive to give a three-dimensional bundle. The use of intermediate flat arrays means that these can be prepared and stored, and custom two-dimensional arrays can be prepared by selecting and attaching together different one-dimensional arrays. This stepwise assembly procedure provides inspection at each step, minimized losses due to errors or low binding efficiency of one rod or tubule, and provides flexibility to assemble new patterns of reactants.

For general clinical use it is important to have identifiers on the slide holding the chip, and identifiers may be integral with the chip itself. Figure 5 illustrated chip 40 with array elements 41, and with a barcode 42 printed along one border to provide identification and orientation. In addition, small concentrations of dyes, usually non-fluorescent, may be incorporated into the polymers from which selected tubes are made such that they present a pattern 43 to 44, for example, of one or more numbers, or one or more letters. It is also

useful to have a few cells or elements which do incorporate fluorescent dyes and which serve to calibrate the fluorescence measurements. It is further feasible to introduce dyes into the contents of selected tubes to additionally identify them. Note that the diagonal line 43-44 further indicates that the horizontal rows of tubes from which the array is assembled, are in the proper order. If tubes in an array are out of alignment giving rise to the loss of one tube or rod in one line, this can be readily observed because the entire pattern will show a misalignment.

The embedding material or adhesive used to hold the tubes in a bundled configuration may be opaque, while the tubes and preferably their contents will conduct light along their length. As a final check on the orientation of array elements, one element at a time at one end of the bundle may be illuminated, and the light detected and related to array position at the other at the other end as shown in Figure 6 where bundle 50 with fibers 51 is illuminated by a cathode ray tube (CRT) 52 generated raster 53 which is focused on the distal end of the bundle by lens 54, and the transmitted light recorded by CCD camera 55. Individual spots 56 yield signals 57 that are detected.

An arrangement for detection using epifluorescence is shown diagrammatically in Figure 7 where chip 60 is illuminated by beam 61 generated by lamp 62, which passes through filter 63 to isolate light of a wavelength optimal for exciting fluorescence. A split-beam prism 64 directs this exciting light toward chip 60. The emitted light passes back through the split-beam prism after which the emitted wavelengths are isolated by filter 65 and detected by CCD camera 66. A variety of different systems for detecting fluorescence patterns on chips have been developed, are commercially available, and are known to those skilled in the arts.

As an alternative method to forming fibers before bundling them, one

may first form the fibers by cleaving them from a larger material. In Figure 8, a sheet of adsorbent material 70 is impregnated with a single ligand or receptor. This may be done by dissolving the compound in a solution and then impregnating a sheet of adsorbent paper (e.g. filter paper). A crosslinking agent may be added to attach the receptor to the cellulose base of the paper or other support. Alternatively, one can crosslink paper pulp to the receptor and then form the sheet of paper or felt. This alternative technique provides a more consistent and uniform distribution but requires greater amount of receptor. Either way, this forms the sheet (70). Many different sheets are prepared, wherein each sheet contains a different receptor.

The sheets are then stacked together (like a book) with adhesive and optionally an inert sheet (not impregnated, preferably black) as a spacer between each sheet of paper. This forms a book (71). One then takes the book to a paper cutter or similar sectioning instrument and a very thin strip (72) is cut which resembles the “ribbon” of Figure 1, object (2). The rest of the process is similar to that shown in Figure 1. Multiple strips (72) from different books are stacked to form a bundle (73) which is then cut transversely to form a microarray (74). An adhesive is preferably added to the ribbons to adhere them. Alternatively, an adhesive may be applied to a solid phase or the end of the bundle and the solid phase adhered to the bundle end before sectioning.

The fiber material is preferably glass, metal, plastic or other polymeric material. For coaxial composite fibers, the dissolvable component may be made of a much wider variety of materials. Each material may be a composite of two or more components. The fibers may act as light pipes or total internal reflection fiber optics to transmit positional alignment and information regarding chemical and biological reactions occurring on the surface. The fiber material is preferably chosen to support attachment of cells and molecules of



interest such as oligonucleotides, peptides and polysaccharides. Hollow fibers may be used to store cells in fresh, frozen or dried condition. Light and electrons emitted directly or indirectly from a reaction or component inside the fiber, particularly a hollow fiber such as a capillary, may be amplified and easily detected when the fiber material is made of glass or other transparent or translucent material. The fiber material may contain a component to react with, detect or convert into another form, the light, electrons or other chemical components emitting from the components or reactions occurring in the fiber. Detection of chemiluminescent reactions in or on the fiber are particularly preferred.

Gelling materials used in the present invention may be selected from a large number of such known materials. Polymers such as agarose, gelatin, collagen, xanthene, carrageenan, alginate, or a thermosetting, thermoplastic, chemosetting or UV polymerizing polymer may be used. Non-polymeric gelling materials including waxes and clays may be used. Hydrogels are particularly preferred when a reaction occurring between the agent of interest and an added substance for interrogation requires an aqueous environment. These may be further modified by using thickeners, gums, hardening and crosslinking agents, plasticizers and various combinations of gelling materials. In general, the gelling material should be sufficiently inert to not interfere with an interaction between the binding component and an analyte.

In the present invention, an agent of interest is extracted into an organic solvent which is miscible with either a thermosetting plastic mixture, or one which is polymerized chemically or by UV or ionizing radiation. This may be done by coating the agents with detergents or other reagents, which will make them soluble under the conditions chosen. The mixture is then extruded into long fibers or cast into fibers. The fibers would be identified by tags on the end

of the fiber or by tags on the rolls carrying the fibers, and/or by incorporating different dyes in them. A barcode may also be printed directly near the end of fibers. Thermoplastic polymers may be used when the embedded product is sufficiently thermostable. Some of the fibers may be differently colored to assist in the localization of specific ligands in the array or to identify the array itself.

The solvent may be miscible in the gelling material or it may be extractable or volatile so as to render a porous final product. Porous products are particularly preferred with solid filament fibers that are self-supporting.

The fibers or their gelling material may also contain a dye or other optical absorber so that only analyte/binding components on the surface of each cell are visualized. Such an improvement reduces the effects of diffusion rates through a gel or porous material that may change with temperature, time, type of carrier liquid, etc. A dye that adsorbs UV or emitted fluorescence will reduce fluorescence from non-surface analyte/binding component reactions.

Different dyes (fluorescent or non-fluorescent) may be incorporated into individual fibers, allowing their location in the two-dimensional array to be confirmed.

The solid filaments or capillary tubes comprising the fibers may be adhered to each other by a variety of techniques. If the components are sufficiently heat stable, the fibers may be sintered together. Alternatively, a number of adhesives are known, including cyanoacrylate adhesives. The space between the fibers may be completely filled by adhesive or a monomer which is polymerized. Thermoplastic and gelling materials may also constitute the adhesive by causing a large number of fibers to be held together in a block. Even inert materials such as Teflon tubes may have their surfaces made reactive with sodium metal in a hydrocarbon solvent to etch the surfaces. Non-

chemicals means such as passing an electrical current through the fibers to fuse them may also be used.

The open ends of the capillaries may be sealed against a flat plate, by pressing a deformable material against the surface, evaporating a plastic (e.g. paralene) on the surface, or by sealing with a chemical such as a thermoplastic or thermosetting plastic material.

There are two basic options for making two-dimensional arrays from these fibers. The first is to make and evaluate ribbons, and then to form a set of ribbons into a long rectangular bar, while the second is to make the bar at the outset, and then all of the fibers together in one step. The former option appears the more advantageous since the ribbons can be evaluated individually before being formed into a complete array. Once the two-dimensional array bar is formed, it can be sectioned using conventional microtomes to form a very large number of slices that can be attached, for example, to glass, metal, or plastic. Alternatively, one may first attach the solid phase material to the end of the bundle before sectioning the bundle. This may be performed by first coating either the end of the fiber bundle or the solid phase with, if necessary, an adhesive such as a cyanoacrylate adhesive or a pre-sectioning or post-sectioning sintering.

Dyed fibers would be visible in such arrays to confirm identifications and orientation. In addition, the fibers can be dyed in such a manner that a visible pattern is formed if the array is correctly made, and the pattern may include a name or a number.

An advantage of the present system is that very large numbers of arrays may be cut, and some fraction of them used for standardization. For example, if a bar 100 cm in length were constructed, and if the bar were cut at 100 micron intervals, then 10,000 arrays would be available. If the sections were

10 microns in thickness, then the number of arrays would be 100,000.

If the individual fibers were 100 microns in diameter, there would be 100 fibers per ribbon, and 10,000 in a bar of fibers with a cross-sectional area of 1 cm square. If there were 330 per ribbon, then the total number-would be 108,900, approximately the number of expressed genes postulated to be present in the human genome.

The present invention is the first array to have such a large number of different cells per unit area on a microarray without the binding agent being covalently attached to the chip. It is preferred for the present invention to have at least 100, more preferably 250, 500, 1,000, 5,000, 10,000, 100,000 or a million or more cells per square centimeter of array. These are much higher concentrations than depositable cells formed by microfluidics in commercial microarrays.

To greatly increase the number of cells per square centimeter beyond even these high numbers, one may prepare a large fiber bundle with relatively large fibers and stretch or draw the bundle. While this makes the individual fibers thinner, it does not affect their basic composition or their orientation with respect to each other and cross-section geometry. This technique has the twin advantages of allowing one to make more microarrays and making them smaller. By using conventional 5 micron porous particles (as in the Example below) and a plastic embedding medium such as a low melting point wax, the result are deformable or ductile fibers which may be drawn to very thin fibers of less than 20 microns in diameter. The field of drawing thermoplastic materials is well known per se. Even if not truly drawable through a die, one can pull or extrude plastic materials between rollers to lengthen and reduce the diameter of the fibers. With optional application of gentle heat, one need only pull the ends of the fiber bundle to generate the same lengthening and reducing

of cross-sectional area. With smaller, porous particles, the fibers may be drawn to even thinner dimensions thereby permitting microarrays of up to at least about 10 billion cells per square centimeter of microarray.

In the field of fiber optics, bundles of optical fibers are heated and drawn into extremely thin optical fibers while retaining their registry within the bundle. Likewise, candy canes and candy with cross-sectional designs are prepared by drawing a large block. Even glass beads used for hundreds of years were also prepared by such techniques.

High concentrations of cells in a microarray have been achieved using photolithography where the molecule of interest is synthesized on the microarray cell. However, the compounds generated by photochemistry are limited. Further, chemically bound compounds interact differently from the same compounds when freely suspended. In a biological system, the active moieties may not be freely available for binding. By contrast, the binding agents of the present invention may be merely entrapped in a matrix, fully retaining their chemical and biological activity.

When using porous particles and immobilizing the molecule of interest inside the porous particle, it may be desirable to retain a suitable fluid inside the pores and use an immiscible embedding medium. In this arrangement, the embedding medium may be incompatible with the molecule of interest or its use in a binding assay, yet still be used. For example an aqueous solution may be used to protect proteins and a low melting point wax used to embed the porous particles.

The known photochemical processes of Fodor et al., *Nature* 364:555-6 (1993); Hacia et al., *Molecular Psychiatry* 3:483-92 (1998); and Fodor et al., *Science* 251:767-773 (1991) prepare short peptides and oligonucleotides covalently bound to the supporting chip. The process of amino acid or

nucleotide synthesis inherently limits the practical length. Synthesis of entire proteins or genes on chips is not practical. Additionally, the secondary, tertiary and quaternary structure of the proteins may be important. By contrast, the present invention permits such.

5 Many different arrays may ultimately be required, and some of these, especially those developed for the identification of infectious agents, may need to be changed at frequent intervals. Further, as new disease-associated alleles are discovered, these will need to be incorporated into new arrays. To fill these requirements and allow changes and additions in arrays, it is important to have individual, stable fiber rolls available, and to have the rolls unambiguously identified. Each roll may be identified by the use of micro-stripes applied at short intervals along the roll. In addition, different tubes may have different colors, and non-fluorescent dyes incorporated into the gels serve as identifiers, or bar coding may be printed on individual fibers.

10  
15 Not only can the chips of the present invention be used to identify infectious agents by identifying characteristic nucleic acid sequences, they can also be used for identifying intact bacteria, mycoplasmas, yeast, nanobacteria, and viruses using arrays of immobilized specific antibodies.

This system may be used for the identification of viruses or other  
20 infectious particles isolated by microbanding tubes, WO99/46047. Thus microbes from biological samples, e.g. serum or plasma, may be concentrated, stained with a fluorescent nucleic acid stain such as TOTO-1 or YOPRO-1, and then allowed to find their matching antibodies on the array. They may then be detected by scanning for fluorescence and identified by position. It is equally a  
25 part of the present invention to immobilize microorganisms or other molecules of interest in the described chips, to use them to localize antibodies from a patient's serum, and to then discover the location of the latter using a

fluorescent anti-human antibody, thus diagnosing the disease which elicited antibody production.

By using the present invention, one avoids the difficulties of individually depositing a different reagent on each cell on a solid phase or synthesizing a  
5 different compound at each cell. The former technique is limited by both the possibilities of spilling and mixing reagents, and by limitations in the accuracy of measurement of small fluid volumes. The latter technique is limited by the types of different compounds that can be synthesized on a solid phase surface.

Both prior art techniques are expensive and require elaborate automated  
10 equipment or tedious labor to produce each array individually. By contrast, the present invention for producing microarrays is technically simple and quick, and the batch size may be in the thousands. The only individual effort required for each microarray is the step of cutting.

Because the bundle is maintained, additional fibers or ribbons may be  
15 added to the bundle as needed before sectioning additional arrays. This allows one to detect and measure newly discovered emerging diseases, new proteins, genes or compounds without recreating a completely new bundle.

This invention may be applied in an alternative fashion in which the bundles are stored at user sites, and the arrays only sliced off as needed. This  
20 arrangement may be useful for research purposes where identical arrays are required over the long term, but only a few are required at any one time.

Another alternative to slicing the bundle and using its sections as separate microarrays is to perform the assay with the end of the bundle directly.

After the assay is performed wherein a first sample could be applied to the cut  
25 cross-sectional surface, and washed off, a detector could image the result. One may then mount the bundle in a microtome device, if the assay were not already so mounted before the assay. A blade could then remove the used surface of

the bundle, exposing a fresh surface for the next assay, which would repeat the same steps again. The bundle could thus be used in one machine for a series of up to 100,000 or more assays performed one after another. This arrangement has certain advantages as optical or electrical detection may be performed through the bundle itself with fiber optic fibers or conductive fibers. The detection system may be continuously attached to the bundle while a more general light or electrical energy applied to the end being used for testing. Specifically note Figure 6 where the testing technology may be adapted to a detection system.

The invention also allows different immobilization technologies, different classes of immobilized agents of interest, different classes of analytes, and different types of detection methodologies to be employed on the same chip.

Since channels are reproducible between plates, the location of each channel or cell may be accurately determined by mechanical means. Reference markings on polished edges or other suitable locations, further identify each cell in the array. Current commercially available computer driven two-dimensional drives of sufficient accuracy are commercially available so that each cell may individually be visualized or tested, or material may be added thereto or withdrawn therefrom.

Cut surfaces of each plate may be polished so that matching plates may be opposed to each other with little possibility of cross leakage. Surface treatment with a material repellant to the fluid to be eventually located inside each cell further reduces cross leakage. For example, fluorinating (Teflonizing) or silanizing agents repel water thereby generating sufficient surface tension to reduce cross leakage of cells filled with an aqueous solution.

After sections have been cut from a bundle, they are generally bound to a



solid backing to provide structural support and ease of handling. The solid backing is typically a sheet of plastic or metal although other materials may be used. The attachment is generally done by a permanent adhesive or heat fusion.

Individual cells in the array may be detected or visualized by scanning the entire array or portions thereof (e.g. one or a few cells) with a charged coupled device (CCD) or by illuminating one or a few channels at a time, such as by a condenser lens and objective lens. The absorbance and emission of light may thus be detected. An optical fiber bundle aligned and registering with the microarray may also be used for optically detecting differences between the cells of the microarray.

Detection may be based on a large number of detectable labels including radioactive, enzyme, luminescent, optically absorbent dye, magnetic, spin-labeled, oxidizers or reducers, chemiluminescence, or indirect labels which interact with a detectable component interacting with the agents of interest in the microarray. The preferred detectable labeling system is based on fluorescence, usually epifluorescence. This requires that the interrogating sample be labeled with one or more fluorescent dyes. The amount of test material required is very small since it would be applied to the arrays as a thin dilute film. Hybridization of nucleic acids would be done under conditions of carefully controlled stringency.

To distinguish selected channels, one may either seal off the selected channels and/or fill them with an easily detectable substance. Different colored inks, dyes and colored materials are particularly well suited as well as detectable components similar to or opposite from the detectable component(s) being detected in other cells. Printing methods with drying inks or plastics, sublimation, solvent containing an ink, or ink-jet printing may be used. The indicia so formed permits better alignment or easily detectable marking when

the array is in use. This permits easy optical alignment.

Once the microarray has been used in a binding assay and the ligands are bound to the receptors, in certain instances it may be useful to provide further identification of the ligand. In certain situations one does not know the entire structure of the ligand from the receptor which specifically binds to it. For example, if the ligand is a cell, a macromolecular complex, or a derivitized molecule with the derivitized portion acting as the ligand, etc., further analysis may be desirable. In this situation, one may elute the ligands from the microarray and collect them for further analysis. For antibody/antigen binding, a pH 2-3 environment or other conditions should strip the ligands. For nucleic acid hybridization, raising the temperature should strip the ligands. A variety of other chemical, physical and electrical techniques for breaking such bonds are known per se.

To enhance specificity to the elution process, an electrode may be placed directly over one or a subset of cells and a current passed through the microarray to release the ligands at that location. The electrode may also be part of a micropipette system to collect the released analytes, see U.S. Pat. No. 5,434,049. Preferably, one uses a porous membrane and applies a current on opposite sides of the membrane.

The method used for analysis of the eluate may be capillary electrophoresis, mass spectroscopy or a second binding assay. Convenient to mass spectroscopy, the microarray itself may be introduced into a laser-desorption system incorporated into a mass spectroscopy system wherein bound molecules are desorbed and analyzed.

Once the analytes have been striped from the microarray, the microarray may be reused. This reuse process has the advantage of being standardized by multiple controls over time.

Additionally, if the receptor is attached to the matrix of the microarray by a cleavable linker, one can isolate the analyte by cleaving the linker. Different cells of the microarray may have different linkers or the same linker and subsequent purification may be needed before additional analysis.

5       The previous methodology for preparation of protein chips requires preparation, use and reuse of large numbers of proteins in solution. Proteins in solution are unstable and deteriorate over time. Even if frozen, repeated use may involve repeated freeze-thaw cycles that denature certain proteins as well.

10       By contrast, immobilized proteins have been shown to be stable over long periods of time.

      The following examples are included for purposes of illustrating certain aspects of the invention and should not be construed as limiting.

#### 15       EXAMPLE 1.       FORMATION AND ANALYSIS OF A MICROARRAY

      Antibodies were prepared by affinity purification by reversible binding to the respective immobilized antigens and subsequently immobilized on particulate supports (Poros G, made by PE Biosystems). An Integral 100Q  
20       biochromatography workstation.

      Each antibody support was made by trapping the antibody on a column of Poros G (commercially available Poros particles pre-coated with protein G, a bacterial protein capable of binding many immunoglobulins by their Fc domain) and subsequently cross-linking the antibody and the protein G with  
25       dimethylpimelimidate (following the PE Biosystems protocol) to covalently immobilize the antibody on the Poros particles. Such antibody columns can be reused (with an acid elution of bound antigen) more than 100 times in a

subtractive mode, and are therefore extremely stable. Each antibody support was characterized to demonstrate specificity for a single antigen.

Antibodies directed against human serum albumin (HSA), transferrin (Tf), and haptoglobin (Hp) were used. A mixture of these three supports was made for use in serum subtraction. A total of three supports were used in tests with: 1) rabbit anti HSA, 2) rabbit anti-human Tf and rabbit anti-human Hp and 3) mixed anti-HSA, Tf and Hp. Unmodified BA Poros (commercially available streptavidin coated Poros), was used as a non-antibody control. Thus, a total of four supports were used.

Poros particles are roughly spherical and highly reticulated (with many internal crevices), having a diameter of approximately 5 microns. Attached proteins are distributed over the internal surfaces as well as the exterior surface of the particle. By embedding the particles in such a medium, a sliceable solid matrix in which the antibody was immobilized and fairly uniformly distributed was created. By exploiting the 3-dimensional nature of the support, a slice containing such particles offers greater capacity (for antibody and thus for antigen binding) than a simple flat surface as used in current microarrays.

Each of the four types of antibody-bearing particles was mixed with an approximately equal volume of 0.75% agarose melted in phosphate-buffered saline (PBS). The agarose for the rabbit anti-HSA beads contained green food coloring to distinguish it. Likewise, the anti-Tf and Hp agarose was colored blue, the mixed anti-HSA, Tf and Hp agarose was colored yellow and the Poros BA containing agarose was white (uncolored). Each melted agarose/bead combination was sucked into a length of one mm diameter plastic tubing of 10 ml length attached to a 1 ml syringe and plunged in ice water. In several minutes the agarose gelled into a jelly-like rod containing approximately 50% Poros beads by volume. The four rods thus obtained (each containing one of

the four bead types above with a different protein coating) were laid into an aluminum channel with more melted agarose to form an array of 2x2 parallel rods embedded in a square cross-section bar of agarose.

After the bar gelled, it was removed from the aluminum channel mold, and transverse sections were prepared by slicing thin slices perpendicular to the axis of the bar (and the filaments) and mounted on a glass slide. These sections revealed a pattern of 4 circular areas (the filaments) containing embedded particulate material (carrying immobilized protein) surrounded by clear embedding matrix of agarose by microscopy. The circular zones of embedded beads were more stable and did not split.

In order to test specific protein binding to the beads in the four circular zones of a section forming the microarray, commercially-available HSA and Tf protein were labeled with fluorescein isothiocyanate (FITC) on Cellite (from Sigma). Cellite is a commercial carrier for insoluble FITC. These proteins were dissolved in about 4 ml of 0.4M sodium bicarbonate buffer (~pH 8.3) and added to the dry FITC on Cellite in the following amounts:

~4.5 mg HSA	30 mg FITC on Cellite
~2.8 mg Tf	18 mg FITC on Cellite
~4.5 mg Serum Protein (20 $\mu$ l)	10 mg FITC on Cellite

The reaction was conducted at room temperature for 30 minutes. The Cellite was removed by centrifugation, and the supernatant protein and unreacted dye placed in a centrifugal protein concentrator, where the protein was washed by repeated dilution and re-concentration in buffer. The fluid was centrifuged to remove the Cellite and supernatant recentrifuged with 4 ml sodium bicarbonate buffer until clear.

Sections of the 4-filament array were laid flat on a glass microscope slide and exposed to a solution of fluorescently labeled HSA. During the exposure

of the section, this protein was expected to interact specifically with the antibodies present on two filaments (round areas on the section): these two filaments were those bearing antibodies to HSA and the mixed anti-HSA, Tf and Hp. Labeled HSA was not expected to interact with the filaments carrying antibodies to Tf alone, or to the filament carrying streptavidin alone.

The section was examined under an epifluorescence microscope equipped with a 500 nm low pass filter and a 510 nm high pass filter for fluorescein fluorescence detection and a 35mm camera.

Prior to extensive washing, all four circular Poros zones showed bright fluorescence, with no discernable differences among them. The fact that the Poros zones showed higher fluorescence than the agarose matrix surrounding the filaments is an indication that the pores of the Poros particles remained unclogged and that the particle-containing zones thus allowed freer diffusion of labeled HSA into the section.

The section was then washed extensively in PBS, and re-examined under the fluorescence microscope. The resulting images, captured on 35mm color slides, demonstrate that after washing, the labeled albumin specifically bound to two filaments and was removed from the other two, thus establishing the ability of the section to specifically detect an individual protein. The two specifically labeled filaments were diagonally opposite one another in the 2x2 array, which was consistent with the diagonally opposite positions of the anti-HSA and mixed anti-HSA, Tf and Hp agarose filaments.

## EXAMPLE 2. MANUFACTURE AND USE OF DIAGNOSTIC ARRAY DETECTING AUTOANTIBODIES TO MITOCHONDRIAL OR LYSOSOMAL PROTEINS.

Suspensions of whole isolated rat and mouse liver mitochondria,

lysosomes, and expressed proteins are suspended or dissolved in an aqueous buffer, at 10 mg/ml concentration, and optionally fixed with glutaraldehyde (1%). 1 ml of each preparation is mixed according to the kit instructions with 20 ml of JB-4 (Polysciences) catalyzed infiltration resin prepared by mixing 20  
5 ml of monomer A containing 0.17 g of catalyst. After complete mixing, 40 mL of monomer B containing 0.17 g catalyst is added with stirring. When completely dissolved, 0.8 g of Accelerator is added, the mixture placed in a syringe and injected into 0.0625 inch internal diameter Teflon tubing under anaerobic conditions. Polymerization occurs at room temperature in  
10 approximately 50 minutes. The ends of the tubes are then heat sealed and stored cold until used, or are immediately extruded for use in preparing a fiber bundle. Bundles are prepared by laying 10 or more fibers in parallel, to make a single-layered array, in an elongated Teflon box. Additional JB-4 resin without protein is then poured in, the box briefly evacuated to remove air bubbles, and  
15 the resin allowed to set. Several such flat arrays may then be stacked in parallel to make a three-dimensional groupings, and the whole grouping further vacuum impregnated to form a three-dimensional bundle. After polymerization, the bundle is cut with a steel or glass microtome knife to give sections 5-20 microns thick, and the sections placed on glass slide. The sections are mounted  
20 using Plastic Mount®, or are dried and mounted with Poly-Mount® (Available from Polysciences).

Tests for autoantibodies are done by placing 0.25 mL of a 1:10 dilution of human serum on each chip and incubating the arrays at 25°C for 20 minutes.

The arrays are then rinsed in phosphate buffered saline four times, and are then  
25 immersed in a solution of goat anti-human globulin conjugated with horseradish peroxidase. After a further 20 minute incubation, the arrays are again washed four times with buffer, and then placed in a solution of 3,3',5,5'-

tetramethylbenzidine in an organic base to which is added a hydrogen peroxide solution (0.02%) in a citric acid buffer. An insoluble blue color indicates the presence of autoantibodies.

### 5 EXAMPLE 3: MANUFACTURE AND USE OF A DIAGNOSTIC ARRAY USING HISTOLOGICAL EMBEDDING SUPPORT.

Arrays are prepared which incorporate fixed infectious particles to be used to detect convalescent antibodies appearing late in the history of an infection. These are important in following sentinel populations to determine  
10 what infectious are occurring.

Immuno-Bed GMA water-miscible embedding medium is made up as directed (Polysciences Inc.), and small batches are mixed with different suspensions of fixed selected viruses (average titer  $10^9$  /ml) or fixed bacterial cells (average  $10^7$  particles /ml). The suspension is placed in a syringe and  
15 forced under pressure into Teflon<sup>®</sup> tubing of 1/16-inch internal diameter, and allowed to polymerize at room temperature. The tubing is pre-treated with metallic sodium in an organic medium to provide a surface, which will adhere to epoxy resins. The polymerized fiber is stored in the coiled Teflon<sup>®</sup> tubing in the cold.

20 The arrays are assembled in bundles using jigs to hold the fibers in parallel array, after which the array is infiltrated with an epoxy resin. The finished bundle, which includes sections of Teflon<sup>®</sup> tubing, is sectioned and the sections mounted on glass slides using an epoxy resin mounting medium. The sections are washed to rehydrate them, and then they are exposed to  
25 convalescent antisera. The chips are then extensively washed and exposed to goat anti-human IgG with the covalently attached fluorescent dye fluorescein. Identification of convalescent antibodies is then done by detecting and



measuring fluorescence using a CCD camera.

#### EXAMPLE 4. MANUFACTURE OF DIAGNOSTIC ARRAY USING SINTERED STRIPS

5 Sintered polystyrene sheets 1/16 inch thick are cut into square-cross-section strips and each one exposed to dilute solutions of one monoclonal antibody to a series of infectious agents including viruses such as rhinoviruses, herpes simplex viruses, influenza virus type A, respiratory syncytial virus, varicella-zoster virus (chickenpox), mycobacterium tuberculosis, cytomegalovirus, Epstein-Barr virus, Hepatitis B Virus (surface antigen and separately core antigen) poliovirus (three strains), and others. The strips are rinsed, dried, and glued together with an acrylonitrile adhesive to form a three-dimensional array that is sectioned to produce arrays 5-100 microns thick. Biological samples containing infectious viruses from individuals with viral diseases are fluorescently stained with the nucleic-acid specific dye YOYO-1 (Molecular Probes) and isolated and concentrated using centrifugal microbanding, see WO99/46047, to concentrate the infectious particles into microliter volumes. The concentrated viruses are applied to the array and are agitated mechanically to move the virus particles over the array for one hour.

20 The array is then washed, excess fluid removed by suction, and illuminated with ultraviolet light at 490 nm. The image is captured with an Apogee CCD camera using a 520 nm filter. Quantitative data is obtained from the processed image using the PMIS image analysis program.

#### EXAMPLE 5. MANUFACTURE AND USE OF DIAGNOSTIC ARRAY HAVING IMMOBILIZED OLIGONUCLEOTIDES:

Polystyrene beads (10-50 microns in diameter) from solid phase oligonucleotide synthesis with oligonucleotides covalently attached are  
5 suspended in buffer and packed into hollow glass fibers of 500 microns internal diameter under hydrostatic pressure initially, and then under air pressure up to 500 psi to expel the supporting liquid. The fiber is then briefly heated under controlled conditions to partially sinter the contents. An array of fibers is then prepared following the methods in the examples above, embedded in a low  
10 viscosity epoxy resin with intermittent vacuum to remove air bubbles, and then allowed to set. The bundle is then sectioned using a diamond saw. The array is used in a flowthrough arrangement as described in US Pat. No. 5,943,767.

#### EXAMPLE 6: MANUFACTURE OF MULTIWELLED PLATES

15 Commercially available glass capillary arrays (GCA)(Galileo) are in the shape of a thin disk having a 2.5 cm X 2.5 cm X 0.5 mm thick dimensions. The GCA has approximately 50% of the area composed of 50  $\mu$  holes or approximately 156,000 holes having a total volume of approximately 0.1 ml. The bottom surface of the GCA is glued to a Teflon® sheet with cyanoacrylate  
20 adhesive (SUPERGLUE).

#### EXAMPLE 7: CLONING AND REPLICA PLATING IN GLASS CAPILLARY ARRAYS

A colony of Streptococcus pyogenes Group A and a colony of Group B  
25 were picked from a plate and mixed together in nutrient agar forming a suspension of these bacterial cells (other microorganisms, animal or plant cells are equally applicable) and are diluted to an approximate concentration of

20,000 cells/ml of culture medium. 0.1 ml of this suspension is applied to the surface of the GCA. This yields about 1 cell per 100 holes to ensure only single cell clones result. The GCA is placed in a sterile petri dish, covered and incubated overnight at 37° C.

5 Two additional sterile GCAs without a Teflon® sheet on the bottom are filled with 0.1 ml heated liquid culture fluid supplemented with 1% agarose, cooled until almost solidified and stacked directly on top of the GCA having cloned bacterial cells so that the holes from each GCA are in register. A top sheet of Teflon® is pressed on tightly and the stack is clamped together. The entire stack is turned upside down and incubated for five minutes at room temperature. The entire stack is turned sideways and incubated overnight at 37° C. The stack is then turned upright, unclamped and individual GCAs are separated. The original GCA is retained for further use.

10 Each of the two added GCAs is placed in a glass flask, attached to a lyophilizer and vacuum dried for 1 hour. The GCAs are removed and 0.1 ml of FITC conjugated antibody to Streptococcus Group A (DIFCO) is added to each GCA and incubated at room temperature for 10 minutes. Each GCA is then blotted on an adsorbent tissue (KIMWIPE) to remove fluid. The microarray is washed by submersion in PBS and blotted dry again. The fluorescent holes in the GCAs and bacteria containing holes in the original GCA are detected using a CCD scanner which gives 12.5  $\mu$  pixels and is capable of a resolution of 25  $\mu$  needed to detect holes which contain cell clones.

15 The scanner is first set to scan for fluorescence and then for absorbance to detect the presence of bacterial clones. Absorbance indicating holes align in both of the two added GCAs. Fluorescent holes align to some but not all of the holes containing bacterial clones in the original GCA.

#### EXAMPLE 8: SELECTING MONOCLONAL ANTIBODIES

Monoclonal antibody-secreting hybridomas in suspension are diluted to approximately 20,000 cells/ml RPMI 1640 + 5% fetal calf serum culture solution and 0.1 ml is added to the GCA of EXAMPLE 6 and the method of EXAMPLE 7 repeated except for incubation being at 37° C in a CO<sub>2</sub> incubator for two days and the GCAs being pretreated with 10% fetal calf serum for 30 minutes. An additional GCA is filled with protein-free saline solution, stacked and clamped. The stack is not turned at all but incubated at room temperature for 15 minutes, unclamped and then vacuum dried as before. 0.1 ml of FITC-conjugated goat anti-mouse immunoglobulin is added to the additional GCA, incubated, removed, washed and scanned for fluorescence as before. Antibody secreting hybridomas are deduced from the location of fluorescence on the GCA.

#### EXAMPLE 9: SCREENING LIBRARIES OF PROTEINS FOR BIOLOGICAL PROPERTIES

Human serum proteins are separated by 2-dimensional electrophoresis as per Baekkeskov et al., Diabetes 38(9): 1133-41 (1989). 200 spots are punched from the gel and the individual proteins dialyzed in 1 ml of PBS. 1 ml of the protein solutions are mixed with 40 mg of acrylamide monomer with catalyst and pumped into 1 mm internal diameter, one meter long polypropylene tubes, the ends heat sealed and each tube tagged. A number of control tubes are prepared with various dyes for easy identification of the correct orientation of the microarray when formed. The acrylamide is allowed to polymerize overnight. The tubes are aligned in a bracket and glued between rows as above. The bundle is cut by a microtome under freezing conditions into 10 micrometer thick slices and the microarray is immediately fixed on a plastic

sheet.

Mouse monoclonal antibodies to the following antigens (Vector Labs) are individually contacted to a separate microarray, incubated, washed, dried and followed by contacting with FITC-conjugated (fluorescein-labeled) goat anti-mouse IgG and scanned as in EXAMPLE 8 above. Insulin, calcitonin, glucagon, epidermal growth factor, interferon, CEA, prostatic acid phosphatase and human IgG are among the antigens tested for. Both hormone levels and tumor antigen levels are determined in a semi-quantitative manner.

#### EXAMPLE 10: RAPID ANTIBIOTIC SENSITIVITY TESTING

Microarrays are prepared in accordance with EXAMPLE 2 except for filling each tube with nutrient agar mixed with various antibiotics in the following configuration. Five two-fold dilutions across the effective spectrum of useful concentrations of the antibiotics, erythromycin, penicillin V, tetracycline, ampicillin, trimethoprim/sulfamethiozole, cefaclor, ofloxacin and nitrofurantoin and 10 two-fold dilutions of 34 new compounds, each a candidate for use as an antibiotic are used.

A colony of an unknown sample of *E. coli* grown from a patient's urine was suspended in 1 ml nutrient broth supplemented with either fluorescein acetate or trypan blue and placed on each of two microarrays and incubated at 37°C. The microarray is scanned for fluorescence and for absorbance at the beginning and after 30 minutes incubation. Microarray cells with detectable increases in fluorescence (scanned fluorescence minus fluorescence from initial scanning) were considered to have growing cells. Microarray cells with increases in trypan blue absorbance from the beginning to 30 minutes were considered to have dead cells. Minimal inhibitory concentrations (MICs) and minimal bactericidal concentrations (MBCs) were thus determined. The

possible effectiveness of the new candidate compounds was likewise deduced.

Another 1 ml of saline containing another suspended colony of the unknown sample of *E. coli* was plated on conventional Mueller-Hinton plates with antibiotic disks and incubated overnight. MICs were determined the next day based on the diameter of the zone of inhibition. The MICs from the microarray are comparable to standardized growth inhibition measurements. For example, for nitrofurantoin, the zone diameter from a 300 mcg disk in millimeters is >17 mm susceptible, 15-16 mm intermediate and <14 mm resistant which corresponds to a MIC in mcg/ml of <32, 64 and >128 respectively. Two-fold dilutions of nitrofurantoin in the microarray are at 16, 32, 64, 128 and 256 mcg/ml.

The method is repeated with known strains of *E. coli* having known differing levels of antibiotic resistance and with many different common microorganisms with different levels of antibiotic resistance. The results indicate which of the 34 candidate compounds are to be tested further as potential antibiotics.

#### EXAMPLE 11: ANTICANCER DIAGNOSTIC AND DRUG SCREENING

Microarrays are prepared according to the method in EXAMPLE 2 with a alkaline-lysed and protease K-digested suspension of various fresh cells from a leukemia patient, several leukemia cell lines (such as, HTB, ATCC), normal peripheral white blood cells and normal bone marrow cells. The microarrays are heat denatured and a digoxigenin-labeled DNA probe for the following genes: N-myc, C-myc, K-ras, p53, HER-2/neu and a candidate DNA probe for diagnostic purposes are applied thereto. Texas Red-labeled anti-digoxigenin antibody is added and the pattern and amount of binding are determined.

## EXAMPLE 12: HEPATITIS TESTING

It is desirable to know the type of viral hepatitis and its stage of infection to best treat a patient. A microarray is prepared as in EXAMPLE 2 except that ten, 2-fold dilutions of mouse monoclonal antibodies to HAV, HBsAg, HBeAg, HCV, HDV and HEV and 2 fold dilutions of the same antigens are used. Three tubes of each are prepared and used in the microarray along with a pattern of controls. Approximately three drops of serum sample is contacted with the microarray, incubated in a 37°C water bath for 10 minutes and washed four times with PBS. 1 ml of a mixed reagent of fluorescein-labeled monoclonal antibodies to non-overlapping epitopes of each of the antigens, fluorescein-labeled mouse anti-human IgG and rhodamine-labeled mouse anti-human IgM is added to the microarray, incubated for 10 minutes in a 37°C water bath and washed four times with PBS. The microarray is scanned for fluorescence at both the wavelength of fluorescein and rhodamine emission and the results determined by which cells of the microarray demonstrate fluorescence and its level.

The microarray is designed for both initial diagnosis and for monitoring treatment and remission by detecting antigens and antibodies in convalescent serum. Two-fold dilutions and measuring the level of fluorescence at each cell provide quantitative results.

## EXAMPLE 13: SCREENING ACTIVE COMPOUND CANDIDATES

Microarrays are prepared according with EXAMPLE 2 except 380 new candidate compounds are introduced into the fibers. Three drops of a solution containing the glutamate receptor 2 are added to the microarray followed by incubation at 37°C for 10 minutes. The microarray is washed and dried as before. A 1:10 dilution of mouse monoclonal antibody to glutamate receptor 2

(Vector Labs) is added, incubated, washed and dried as before. FITC-conjugated goat anti-mouse IgG is added and the microarray scanned.

Fluorescent cells correspond to compounds that bind to the receptor. Since this receptor is involved in learning, memory, seizures and other neurological conditions by binding the neurotransmitter glutamate, both agonists and antagonists are of pharmacological interest.

It will be understood that various modifications may be made to the embodiments disclosed herein. Therefore, the above description should not be construed as limiting, but merely as exemplifications of preferred embodiments. Those skilled in the art will envision other modifications within the scope and spirit of the claims appended hereto.

All patents and references cited herein are explicitly incorporated by reference in their entirety.